

CRF Problem Report

The Scientific and Technical Information Center (STIC) experienced a problem when processing the following computer readable form (CRF):

Application Serial Number: 09/6/49/3 (1907)				
Filing Date:				
Date Processed by STIC: $2-5-00$				
STIC Contact: Mark Spencer, 703-308-4212				
Nature of Problem:				
The CRF (was):				
(circle one) Damaged or Unreadable (for Unreadable, see attached)				
Blank (no files on CRF) (see attached)				
Empty file (filename present, but no bytes in file) (see attached)				
Virus-infected. Virus name: The STIC will not process the CRF				
Not saved in ASCII text				
Sequence Listing was embedded in the file. According to Sequence Rules, submitted file should only be the Sequence Listing.				
Did not contain a Sequence Listing. (see attached sample)				
1 Other: Non-compliant sequence format.				

PLEASE USE THE CHECKER VERSION 3.0 PROGRAM TO REDUCE ERRORS. SEE BELOW FOR DETAILS:

Checker Version 3.0

The Checker Version 3.0 application is a state-of the-art Windows based software program employing a logical and intuitive user-interface to check whether a sequence listing is in compliance with format and content rules. Checker Version 3.0 works for sequence listings generated for the original version of 37 CFR §§1.821 – 1.825 effective October 1, 1990 (old rules) and the revised version (new rules) effective July 1, 1998 as well as World Intellectual Property Organization (WIPO) Standard ST.25.

Checker Version 3.0 replaces the previous DOS-based version of Checker, and is Y2K-compliant. Checker allows public users to check sequence listings in Computer Readable form (CRF) before submitting them to the United States Patent and Trademark Office (USPTO). Use of Checker prior to filing the sequence listing is expected to result in fewer errored sequence listings, thus saving time and money.

Checker Version 3.0 can be down loaded from the USPTO website at the following address: http://www.uspto.gov/web/offices/pac/checker

Corrected Diskette Needed

SEQUENCE LISTING

COMMON FOR ALL SEQUENCES. SEQUENCE TYPE: Peptide SEQUENCE UNIT: Amino Acid

TOPOLOGY: Linear

Non-compliant sequence listing format. See attached example of correct format and explaination sheet.

SEQUENCE ID NO: 1

SEQUENCE LENGTH: 19 amino acids

NVPGHERMGRGRTSSKELA

NVPGHERMGRGRTSSKELA

1 5 10 15

SEQUENCE ID NO: 2

SEQUENCE LENGTH: 27 amino acids

RLEAKHRENVPGHERMGRGRTSSKELA

1 5 10 15 20 25

SEQUENCE ID NO: 3

SEQUENCE LENGTH: 17 amino acids

RLEAKHRENVPGHERMG

1 5 10 15

SEQUENCE ID NO: 4

SEQUENCE LENGTH: 12 amino acids

MGRGRTSSKELA

1 5 10

SEQUENCE ID NO: 5

SEQUENCE LENGTH: 15 amino acids

Appendix A To Subpart G to Part 1-Sample Sequence Listing

<110> Smith, John

and the first state of the second

Smith, Jane

<120> Example of a Sequence Listing

<130> 01-00001

<140> US 08/999,999

<141> 1998-02-28

<150> EP 91000000

<151> 1997-12-31

<160> 2

Federal Register

<170> PatentIn ver. 2.0

<210> 1

<211> 403

<212> DNA

<213> Paramecium aurelia

<220>

<221> CDS

<222> 341..394

<300>

<301> Doe, Richard

<302> Isolation and Characterization of a Gene Encoding a

Protease from Paramecium sp.

<303> Journal of Fictional Genes

<304> 1

<305> 4

<306> 1 - 7

<307> 1988-06-20

<400> 1

ctactctact ctactctcat ctactatctt ctttggatct ctgagtctgc ctgagtggta 60

ctcttgagtc ctggagatct ctcctctcac atgtgatcgt cgagactgac cgatagatcg 120

ctgactgact ctgagatagt cgageccgta cgagacccgt cgagggtgac agagagtggg 180

cgcgtgcgcg cagagcgccg cgccggtgcg cgcgcgagtg cgcggtgggc cgcgcgaggg 240

ctttcgcggc agcggcggcg ctttccggcg cgcgcccgtc cgcccctaga cctgagaggt 300

cttctcttcc ctcctcttca ctagagaggt ctatatatac atg gtt tca atg ttc

Met Val Ser Met Phe

age ttg tet tte aaa tgg cet gga ttt tgt ttg ttt gtt tgtttgete

Ser Leu Ser Phe Lys Trp Pro Gly Phe Cys Leu Phe Val

10

15

<210> 2

<211> 18

<212> PRT

<213> Paramecium aurelia

<400> 2

3; No.; 104/Monday, June 1, 1998/Ru.

29643

Met Val Ser Met Phe Ser Leu Ser Phe Lys Trp Pro Gly Phe Cys Leu

1

5

10

Phe Val

× . 1 ·	-tifior Library	Definition	Comments and format	Mandatory (M) or optional (O)
	The state of the s	Berline Lander	A STATE OF THE STA	
3.	<110>	Applicant	Preferably max. of 10 names; one name per- preferable format: Surname, Other Names a or Initials.	17777
•	<120>	Title of levention	Or mulais.	M. Samuelore
•	<130>	File Reference	Personal file reference	M when filed prior to assignment of appl. number.
-	<140>	Current Application Number,	Specily as: US 07/999,999 or PCT/US96/99999-	M, iLavailable.
	<141>	l	Specify as: yyyy-mm-dd	M, if available.
	<150>	Prior Application Num-	Specify as: US 07/999,999 or PCT/US96/99999	M, if applicable include priority documents under 35:USC 119 and 120.
	<151>	Prior Application Filing Date.	Specify as: yyyy-mm-dd	M, if applicable.
	<160> <170>	Number of SEQ ID NOs Software	Count includes total number of SEQ ID NOs Name of software used to create the Sequence Listing.	0
	<pre><210></pre>	SEQ ID NO:#:	Response shall be an integer representing the SEQ-ID NO shown.	M
	<211>	Length	Respond with an integer expressing the number of bases or amino acid residues.	М.
	Numeric Iden-	Definition	Comments and format	Mandatory (M) or optional (O).
	<212>	Туре	Whether presented sequence molecule is DNA, RNA, or PRT (protein). If a nucleotide se-	M.
			quence contains both DNA and RNA trag-	, j
			ments, the type shall be "DNA." In addition, the combined DNA/RNA molecule shall be further described in the <220> to <223> feature	And the second s
	<u> </u>	Organism	section. Scientific name, i.e. Genus/ species, Unknown or	M
			Artificial Sequence. In addition, the "Unknown" or "Artificial Sequence" organisms shall be further described in the <220> to <223> feature	• • • • • • • • • • • • • • • • • • •
	<220>	Feature	section. Leave blank after <220>. <221-223> provide for a description of points of biological significance	M, under the following conditions: if "n," "Xaa," or a modified or unusual L-amino acid or modi-
			in the sequence	fied base was used in a sequence; if ORGA- NISM is "Artificial Sequence" or "Unknown"; if molecule is combined DNA/RNA"
	<221>	Name/Key	Provide appropriate Identifier for feature, preferably from WIPO Standard ST.25 (1998), Appendix 2, Tables 5 and 6.	M, under the following conditions: if "n," "Xaa," or a modified or unusual L-amino acid or modified base was used in a sequence.
	<222>	Location	Specify location within sequence; where appro- priate state number of first and last bases/ amino acids in feature.	M, under the following conditions: if "n," "Xaa," or a modified or unusual L-amino acid or modified base was used in a sequence.
	<223>	Other Information	Other relevant information; four lines maximum	M, under the following conditions: if "n," "Xaa," or a modified or unusual L-amino acid or modi- fied base was used in a sequence; if ORGA- NISM is "Artificial Sequence" or "Unknown"; if molecule is combined DNA/RNA.
	<300>	Publication Information	Leave blank after <300>	0.
	<301>		Preferably max of ten named authors of publica- tion; specify one name per line; preferable for- mat: Surname, Other Names and/or Initials.	O. ··
	<302>	Title		O
	<302>	Journal		0.
	<304>	Volume		O. O.
	<305>	1 -		0.
	<306>	Pages	Journal date on which data published; specify as	Ŏ.
•	<307>		yyyy-mm-dd, MMM-yyyy or Season-yyyy. Accession number assigned by database include	O.
		Number.	ion database name.	0 .
•	<309>	Database Entry Date	Date of entry in database; specify as yyyy-mm- dd or MMM-yyyy.	O
	<310>	Patent Document Number.	Document number; for patent-type citations only. Specify as, for example, US 07/999,999. Document filing date, for patent-type citations	O
	<311>		only; specify as yyyy-mm-dd. Document publication date, for patent-type cita-	0.
	<312>	r dolication Date	tions only: specify as yyyy-mm-dd.	
	<313>	Relevant Residuesz	FROM (nosition) TO (position)	0.
	<400>		SEQ ID NO should follow the numeric identifier and should appear on the line preceding the actual sequence.	м.
			adual sequence.	